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depot of a composition containing said azalide antibiotic on the eye.

Claim 47. (Amended) The process according to claim 45, wherein said azalide antibiotic is a compound of formula (I):

Jube C2

wherein  $R^1$  and  $R^2$  each independently represent a hydrogen atom or methyl group.

Claim 46. (Amended) The process according to claim 41, wherein said azalide antibiotic is azithromycin.

Claim 36. (Amended) The process according to claim 48, wherein said applying provides a therapeutically effective concentration of azalide antibiotic within a tissue of the eye for at least, 8 hours.

Claim 50. (Amended) The process according to claim 49. wherein said applying provides a therapeutically effective concentration of azalide antibiotic within a tissue of the eye for at least 12 hours.

Claim 51. (Amended) The process according to claim 50, wherein said applying provides a therapeutically effective

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 $\lambda$ concentration of azalide antibiotic within a tissue of the eye for at least 18 hours.

Claim 53.8 (Amended) The process according to claim 45, wherein said depot is an aqueous polymeric suspension of said azalide antibiotic.

Claim 156. (Amended) The process according to claim 156. wherein said depot is a composition selected from the group consisting of an aqueous suspensions, ointments, and inserts.

Claim 156. (Amended) The process according to claim 15. wherein said depot remains for at least 30 minutes after administration.

Claim 61. (Amended) A topical ophthalmic composition comprising an aqueous polymeric suspension comprising water, 0.01% to 1.0% of an abated antibiotic and 0.1 to 10% of a polymeric suspending agent.

Claim 62. (Amended) The composition according to claim 61, wherein said suspension further comprises an additional medicament selected from the group consisting of antibiotics, antivirals, antifungals, anesthetics, anti-inflammatory agents, and anti-allergic agents.

Claim 3. (Amended) The composition according to claim 52, wherein said additional medicament is contained in the amount of from 0.01 to 5.0%.

Claim 64. (Amended) The composition according to claim 62, wherein said additional medicament is selected from the group consisting of amikacin, gentamycin, tobramycin, streptomycin, netilmycin, kanamycin, ciprofloxacin, norfloxacin, ofloxacin, trovafloxacin, lomefloxacin, levofloxacin, enoxacin, sulfonamides, polymyxin, chloramphenicol, neomycin,

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paramomomycin, colistimethate, bacitran, vancomycin, tetracyclines, rifampins, cycloserine, beta-lactams, cephalosporins, amphotericins, fluconazole, flucytosine, matamycin, miconazole, ketoconazole, corticosteroids, diclofenac, flurbiprofen, ketorolac, suprofen, lodoxamide, levocabastin, naphazoling, antazoline, and pheniramimane.

Claim 65. (Amended) A topical ophthalmic composition comprising an effective amount of an azalide antibiotic, an ophthalmically acceptable carrier, and an additional medicament selected from the group population of antibiotics, antivirals, antifungals, anesthetics, anti-inflammatory agents, and anti-allergic agents.

Claim 6. (Amended) The composition according to claim 65, wherein said azalide antibiotic is azithromycin.

Claim 25. (Amended) The composition according to claim 25.20 wherein said composition is fluid; said azalide antibiotic is contained in an amount of from about 0.01 to 2.0%; and said additional medicament is contained in an amount of from about 0.01 to 5.0%.

Claim 73. (Amended) The process according to claim 70, wherein said topically applying comprises supplying a depot of a composition containing said azalide antibiotic on the eye.

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Claim 83. (Amended) The process according to claim 70, wherein said azalide antibiotic is a compound of formula (I):

mbs CS

wherein  $R^1$  and  $R^2$  each independently represent a hydrogen atom or methyl group.  $u_0$ 

Claim 84. (Amended) The process according to claim 83, wherein said azalide antibiotic is azithromycin.

Claim 85. (Amended) The process according to claim 70, wherein said applying provides a therapeutically effective concentration of azalide state of the eye for at least 8 hours.

Claim 6. (Amended) The process according to claim 86, wherein said applying provides a therapeutically effective concentration of azalide antibiotic within a tissue of the eye for at least 12 hours.

Claim 87. (Amended) The process according to claim 26, wherein said applying provides a therapeutically effective concentration of azalide antibiotic within a tissue of the eye for at least 18 hours.

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Claim 7. (Amended) The composition according to claim 65, wherein said composition is fluid; said azalide antibiotic is contained in an amount from at least about 5.0%, and said additional medicament is contained in an amount of from about 0.01 to 5.0%.

Claim 98. (Amended) The composition according to claim 55, wherein said composition is fluid; said azalide antibiotic is contained in an amount from about 0.1 to about 5.0%, and said additional medicament is contained in an amount of from about 0.001 to 5.0%.

Claim 101. (Amended) A topical ophthalmic composition comprising an aqueous polymeric suspension comprising water, at least about 5.0% of an azalide antibiotic, and 0.1 to 10% of a polymeric suspending agent.

Claim 102. (Amended) A topical ophthalmic composition comprising an aqueous polymeric suspension comprising water, from about 0.1 to about 650% of an azalide antibiotic, and 0.1 to 10% of a polymeric suspending agent.

### REMARKS

Support for the amendment to Claim 45 can be found, inter alia, in cancelled Claim 52; support for the amendment to Claim 61 can be found, inter alia, in Claim 65; and support for the amendment to Claim 65 can be found, inter alia, in cancelled Claim 67. The remaining amendments are relatively minor, and primarily editorial in nature.

On pages 2-3 of the Office Action, the Examiner rejects Claims 45-60, 70-87 and 95-96 under the doctrine of